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Nathalie Maitre demonstrates the "puffer system" in the VKC Psychophysiology Lab.

Tapping the Sensory System's Power BY JAN ROSEMERGY

N athalie Maitre is passionate about babies, especially infants cared for in neonatal intensive care units (NICUs). An assistant professor of Pediatrics in Neonatology and adjunct in Physical Medicine and Rehabilitation, Maitre, Ph.D., M.D., is a clinician and translational researcher set on discovering interventions to help these infants flourish.

Maitre corrects the misperception that NICUs provide care only for premature infants. "We also care for babies with congenital abnormalities, cardiac problems, birth asphyxias or trauma, infections at birth, and more. Together, such infants contribute greater than 60% of children with cerebral palsy, vision and hearing problems, and intellectual disabilities."

Maitre's research focuses on early identification and rehabilitation of neurodevelopmental problems in high-risk infants. "My clinical duties allow me a unique perspective on care and rehabilitation and have influenced my focus on interventions that take advantage of neural plasticity in the early years," she said.

Maitre focuses on the sensory system because it is the basis for learning in infancy. Brain processes involve not only developing specific senses like vision or hearing but connecting senses into networks. The development of these neural networks is the basis for more complex processes and higher order cognitive functions. "If you can make a difference at the beginning, in the first months and years of life, then hopefully it amplifies later in life," Maitre said.

Among Maitre's research collaborators is Sasha Key, Ph.D., associate professor of Hearing & Speech Sciences and VKC Psychophysiology Lab director. They are developing objective measures of neurophysiologic function in infants to validate early therapeutic interventions.

Maitre thinks of the brain less as a computer and more as a changing internet. The basic principle underlying her work, like most of her colleagues in this field, is that neural stimulation plus behavioral training can rewire the brain and the peripheral nervous system.

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Known for My Strengths Rethinking Our Introductions BY COURTNEY TAYLOR AND ERIK CARTER

D isability is a *natural part of the human experience and in no way diminishes the right of individuals to participate in or contribute to society.* This simple statement is prominent within almost every piece of federal legislation focusing on people with intellectual and developmental disabilities. Being part of a community is at the heart of what it means to flourish as a human. Yet the experiences and relationships that make up everyday community life remain elusive for far too many people with significant disabilities.

Although much progress has been made over the last few decades, many employers, community groups, congregational leaders, and others still struggle to welcome and weave people with disabilities into the activities and relationships that are part of life in community. They may have difficulty imagining a meaningful place for someone described as having Down syndrome or autism. They may be uncertain of what someone with a label of fragile X syndrome or multiple disabilities has to bring to their business, civic organization, synagogue, or neighborhood group.

Think about how we typically talk about disabilities. It is often in terms of what someone *cannot* do or struggles to do. Such a message has inadvertent consequences. For example, when an employer considers, "What could a person with an intellectual disability contribute to this business?" or a youth ministry leader wonders, "How do we include a teenager with autism in our programs?" — all they might have is an image of what a person with that particular label cannot do. That makes for a challenging introduction. It is difficult to envision a place in a community for someone only on the basis of a list of deficits. How we introduce people surely matters.

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Director's Message Compassion—For Others and for Ourselves

he lure of a mother's lullaby in a Neonatal Intensive Care Unit. Families and Ambassadores

cheering as Next Steps at Vanderbilt students graduate. Wiping away tears while listening as families affected by autism share their stories. A mentor speaking words of encouragement to a postdoctoral fellow about to lead a professional symposium for the first time. LEND and UCEDD trainees sharing transformational leadership experiences from the past year. These are but a few instances of compassion that abound in the lives of Vanderbilt

Kennedy Center researchers, clinicians, professional staff, trainees, and students.

Everyday, all of us in our VKC community work to make positive differences, directly or indirectly, in

the lives of children and adults with developmental disabilities and their families. We are focused on

giving support. To make that possible, we also support one another.

We support young student and postdoctoral researchers by engaging them in labs led by senior researchers, by featuring their research at VKC Science Day, by providing VKC Travel Awards to present their research at scientific conferences.

We support young and seasoned researchers alike by partnering with donors to provide Nicholas Hobbs

Discovery Grants to seed innovative research. We support journal clubs that bring together faculty, staff, students, and community members to learn and brainstorm. In "More About That," program staff experience the broad range of our Center's work. In "Developmental Disabilities Grand Rounds," we hear the work of our own researchers and discover new possibilities for collaboration.

As I write, we are working on grant progress reports, submitting grant applications, recruiting for research studies, concluding semester exams and submitting grades, doing annual performance evaluations, and more. In the middle of these activities, as a community, we are grieving the loss of a long-time staff member and are comforting one another.

As we care for others, we also must care of ourselves, whoever we are, whatever we do. Each of us can learn and practice ways to take better care of ourselves by living attentively and finding things for which we can be grateful. Today we are grateful for this community.

Passionate About Developmental Disabilities Research 2013 Gatlinburg Conference BY ELIZABETH TURNER

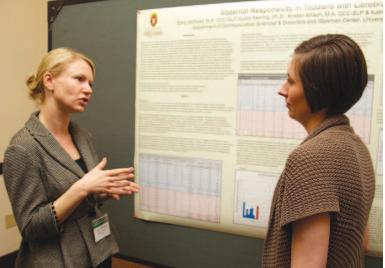
More than 200 students and professionals made their way to San Antonio, Texas, March 6-8, for the 46th Annual Gatlinburg Conference on Research and Theory in Intellectual and Developmental Disabilities. This year's theme was "Newborn Screening for Intellectual and Developmental Disabilities."

"The development and expansion of newborn screening has been enormously important in the prevention of developmental disabilities," said Elisabeth Dykens, Ph.D., VKC director and Conference Chair.

The Gatlinburg Conference is the preeminent national conference on intellectual and developmental disabilities where both young and seasoned researchers, from graduate students to senior faculty, come together for vigorous dialogue.

"This was my first time attending this conference, and I was thrilled and honored that my proposal for a symposium on *Infant Sensory Processing and Neurodevelopmental Outcomes* was selected," said Nathalie Maitre, Ph.D., M.D. (VKC/Vanderbilt U). "Professionally, it was the first time I was given the opportunity to organize the content and structure of research to make it more meaningful and accessible to a diverse audience of scientists, advocates, and clinicians. It is a rare and wonderful opportunity to be learning and communicating with the few people in this country who feel just as passionately about disabilities research as we do!"

Vanderbilt participants included 12 faculty, 16 graduate students and postdoctoral students, 2 research staff members, and 4 VKC staff who coordinated the conference. This year's conference was the third in the VKC's 5-year coordination commitment with the *Eunice Kennedy Shriver* National Institute of Child Health and Human Development.



A Gatlinburg Conference attendee presents her findings during one of three poster sessions.

Between the 4 plenary sessions were 12 symposia chaired and presented by professionals from universities around the continent. Topics ranged from mindfulness-based stress reduction to Rett syndrome to social vulnerability of those with intellectual and developmental disabilities.

Marisa Fisher, VKC postdoctoral fellow who received a Theodore Tjossem Postdoctoral Award to attend the Conference, chaired a symposium on social vulnerability. "I have attended the Gatlinburg Conference every year since I entered graduate school," Fisher said. "As a postdoc, I was eager to chair a symposium. What came together was a wonderful group of researchers interested in

> the same topic. Our symposium was a success. Not only was it well attended, but also it generated more research questions and the promise of future collaborations with new colleagues."

NICHD staff provided updates and a panel discussion. Questions asked included the possible effects of sequestration and the future of the grant application/renewal process.

In all, 21 travel awards were given to undergraduates, graduate students, and postdoctoral fellows, many of whom attended for the first time.

Symposium and poster abstracts are available at kc.vanderbilt.edu/gatlinburg.

The 47th Annual Gatlinburg Conference will be held March 5-7, 2014, at the Hotel Allegro in Chicago. The theme will be "The Co-Occurrence of Mental Illness and Developmental Disability."



Elisabeth Dykens

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TAPPING THE SENSORY SYSTEM'S POWER from page 1

Musical Pacifiers

It is common practice in NICUs to use special pacifiers to encourage infants to suck and thus learn

to feed. Maitre's idea was to harness the power of a baby's mom singing a lullaby—an auditory stimulus— to strengthen neural connections among sucking, breathing, and swallowing. The lullabies sung were modified by a music therapist, Olena Chorna, MT, to fit patterns that an infant's auditory system would recognize and respond to. "While the infants are sucking, they're learning there's a right rhythm and pressure, and that when they suck the right way, they get their mother's voice singing, a positive stimulus for their developing brains."

With support from a VKC Hobbs Discovery Grant, Maitre and Chorna conducted a randomized clinical trial of

this pacifier-activated lullaby and measured how infants fed. After just 5 days, the infants with the musical pacifiers tripled their suck rate. They increased their number of feeds per day by three compared to baseline, while the comparison group increased their number of feeds per day by one. "Babies have eight feeds a day, so an increase of three



is huge," Maitre said. The infants with the musical pacifiers increased their feeding volumes, they had fewer days of tube feeding, and had decreased length of hospital stay.

Puffer System

Maitre and Key have developed a multisensory paradigm to measure the perception of light touch

> simultaneous with perception of a speech sound, which permits insight into touch and sound not as separate modalities but as the two combined. Wearing an ERP sensory "hat" to measure brain electrical activity and with a finger in a special mold, a seated child is distracted by a muted video while experiencing air puffs or speech sounds or simultaneous air puffs and speech sounds. They have found a significant difference between how touch is perceived alone and how touch is perceived in combination with sound.

"This work has treatment implications for developmental disorders like autism in which tactile or multisensory processing is affected," Maitre said.

Maitre values the collegiality of

Vanderbilt and the VKC. "I don't feel lonely in how much I care about kids with disability anymore. It's wonderful to belong to a community who also cares so deeply about them."

KNOWN FOR MY STRENGTHS from page 1

Study of Strengths and Flourishing

As part of the VKC UCEDD *Disabilities, Religion, and Spirituality Program*, we are in the midst of a project focused on reshaping such introductions. We have been conducting a statewide study focused on faith and flourishing in the lives of youth and young adults with intellectual disabilities and autism in Tennessee. One of the study's aims is to identify the strengths these young people might have to share with others in their community. The study is funded by the **Martin McCoy-Jespersen Discovery Grant in Positive Psychology**. Martin is remembered by his family and friends as "open, accepting, present," whose "great purpose was to live a happy life" connected to others.

To identify strengths, we asked more than 450 parents to complete a short scale focused on enviable qualities of these youth and young adults. The Assessment Scale for Positive Character Traits (Woodard, 2009) includes 26 different statements, each addressing the extent to which their child shows characteristics like kindness, humor, gratitude, empathy, optimism, forgiveness, and courage.

Gathering this type of "positive" data has implications for fostering inclusion and community

participation. What if we introduced young people to others by their strengths and gifts? Can we think of young people with intellectual and developmental disabilities in this way? Findings from our study to date suggest their parents can see many of these assets.

Drawing from the hundreds of parents and caregivers who participated, consider this small sampling of findings:

- 94% of parents described their child as *happy*.
- 86% say their child had a great sense of *humor*.
- 85% of these young people were described as *thoughtful and helpful to others*.
- 85% said their child shows kindness to others.
- 85% of parents said their child *is thankful for life's simple pleasures*.
- 75% of parents described their child as *courageous*.
- 70% said their child *keeps on trying even when things get hard.*

How many businesses would benefit from hiring someone with qualities like honesty, persistence, and optimism? How many faith communities can find a place for someone known for her gratitude, empathy, and kindness? How many neighbors would be eager to develop a friendship with someone who is funny, happy, and thoughtful?

Does describing people in light of their strengths make any difference? Consider these two introductions: (1) Meet John. He has an intellectual disability, he can't really read, and he is going to need lots of help to get around. (2) Meet John. He has an incredible sense of humor, he loves meeting new people, and he enjoys learning new things. By the way, he has an unforgettable bear hug. Which John would you be more eager to meet?

Youth and young people with disabilities have wonderful strengths and gifts to share. Too often those potential contributions get overlooked. As we strive to equip communities to more meaningfully include people with disabilities, let's not overlook the introduction. Leaders and members of all our communities must see people first in terms of the gifts they have to bring.

Information from this study will be made available through a practical guide. To receive a forthcoming practical guide, email courtney.taylor@vanderbilt.edu. Courtney Taylor. M.Div., is VKC associate director of Communications and Dissemination. Erik Carter, Ph.D., is associate professor of Special Education.

VANDERBILT KENNEDY CENTER FOR EXCELLENCE IN DEVELOPMENTAL DISABILITIES



Pathfinder staff pictured left to right: Cecilia Melo-Romie, Tracy Beard, Lydia Wingo Kane, Carole Moore-Slater, Aram Torabiam, Megan Hart, Alexander Santana, and Carolina Meyerson.

Pathfinder's Path BY COURTNEY TAYLOR

ver the course of the last year, the staff at Tennessee Disability Pathfinder has provided information and referral assistance to over 2,000 callers from 87 of Tennessee's 95 counties, 24 states, and 34 countries of origin. In addition to HELPline assistance, an analysis of the Pathfinder Web site indicates it had more than 54,800 visitors from 191 cities and towns across the state. These impressive numbers are a far cry from Pathfinder's humble beginnings. With a current staff of 8, plus interns and Work Study students, Program Director Carole Moore-Slater, M.S., who recently announced she will retire in June, remembers when it was just herself and a part-time administrative assistant.

"At first, Pathfinder was just a toll-free helpline with a couple of people in the office," said Moore-Slater. The program began with a grant written by VKC Director of Communications Jan Rosemergy, Ph.D., funded by the Tennessee Council on Developmental Disabilities.

"The first year I think we maybe had a couple hundred calls, mainly from folks in Middle Tennessee," Moore-Slater said. "Our reach certainly wasn't statewide and beyond like it is today, but we learned so much in that first year. We grew by gathering information about services in each county. We built a database and a Web site to house all that information. Then, we realized that to become a clearinghouse for disability-related resources and services, we had to get out from behind our desks and get into the communities. So, we began to network and build relationships to learn first-hand about what services were available."

Outreach in the community paid off and eventually, the Pathfinder database grew to include over 2,100 agencies. Pathfinder's programming also began to expand. Its growing staff began conducting community trainings on topics such as recreation and leisure, health care, and disability etiquette.

Access Nashville, a service-learning project to assess the accessibility of local restaurants, began and would eventually train over 800 students and survey 500 restaurants, all of which are listed on the Nashville Convention and Visitors Bureau's Web site.

Pathfinder also expanded its reach by hiring multilingual staff members, adding a database (Camino Seguro) of agencies that have a Spanishspeaking staff member, starting a support group

for Spanish-speaking families, and holding annual cultural diversity conferences to address issues relevant to the growing number of immigrants and refugees who have family members with disabilities from diverse cultural backgrounds now living in Tennessee.

"The Multicultural Outreach Program is an incredible achievement," said Moore-Slater. "What Pathfinder has done to improve services for immigrants

and refugees living in Tennessee who have family members with disabilities is incredible. The support groups alone have contributed so much to the

A family served in a Multicultural

Outreach Program Support group

quality of life of the members. Our staff also has been instrumental in developing the Tennessee Multicultural Alliance on Disability, which is doing such great work with the help of agencies across the state to create better opportunities for immigrants and refugees. I am going to miss being a part of this."

"Carole sees the important value of training through Pathfinder for the future generation of leaders in the disability field," said Elise McMillan, J.D., Pathfinder faculty director and VKC UCEDD co-director. "Students from across the University have benefitted from working as Pathfinder trainees."

Before taking on a leadership role 16 years ago and helping to build Pathfinder into what it is today, Moore-Slater worked as a social worker, as a parent support trainer, and as a teacher at the Susan Gray School. Looking back, she says one major common element of those positions was trying to find information about community services for the children and families, and the difficulty she had in finding them. In part, that is what drew her to Pathfinder's mission.

"When I saw the position open, I was really excited about it," said Moore-Slater. "Trying to streamline a path to our fragmented services in Tennessee was such an important endeavor. I knew it would be the perfect path for me and I have never looked back."

"We are very thankful for Carole's leadership over the last 16 years," said Wanda Willis, executive director of the Tennessee Council on Developmental Disabilities. "She has been the driving force behind Pathfinder. Carole was hired as the first director and from that moment led with vision, creativity, and a fierce commitment to connecting Tennesseans with disabilities, their families, and supporters to information and resources. Pathfinder has been

> heralded nationally and replicated by other states, and a lot of that is due to Carole's leadership. She is someone I personally admire and salute for her accomplishments. She will be missed, but leaves behind a great legacy."

Tennessee Disability Pathfinder is a joint program of the VKC UCEDD and the Tennessee Council on Developmental Disabilities. Funding also is provided by the

Tennessee Department of Intellectual and Developmental Disabilities and the Tennessee Department of Education.

New TRIAD Leadership Roles BY JAN ROSEMERGY

T RIAD staff are assuming new leadership roles, announced Zachary Warren, Ph.D., TRIAD director and associate professor of Pediatrics, Psychiatry, and Special Education.

Pablo Juárez, M.Ed., BCBA, has been named administrative director of TRIAD Professional Development and Training. TRIAD's team provides educational training and consultation to teachers, psychologists, administrative staff, and school districts throughout the state and region. This includes a contract with the Tennessee Department of

Education to train educational staff in evidencebased educational practices for children with autism and related disabilities.

LaTamara Garrett, B.A., has been named the

program coordinator for Early Childhood Training, which includes contracts with the Tennessee Department of Education to conduct



L to R: Pablo Juárez, LaTamara Garrett, and Whitney Loring

regional training for early childhood educators. Garrett is focused on expanding the impact of such training through novel methods and systems, including potentially realizing model classrooms and distance modalities for training.

Whitney Loring, Psy.D., is now program coordinator of Families First, a free training series

> for parents of young children with autism. By May 2013, Families First will have served more than 2,500 family members across the region since the workshops began in 2008.

"With the ever-growing need for autism services, we need training leaders who can translate cutting-edge science into sustainable educational practice," says Warren.

"We are very fortunate to have leaders who understand this critical need and have the vision to realize training programs of true meaning and impact."

Testing Toddler Interventions BY JAN ROSEMERGY

A lthough evidence supports the importance of early intervention for young children with autism spectrum disorders (ASD), more research is needed to identify which specific treatments are most helpful for which children and at what level of intensity. Now underway is a national multisite randomized clinical trial of early intervention for

ASD to provide information on what effects the style and the intensity of treatment (number of hours per week) have on children's development. Led by Sally Rogers, Ph.D. (U California Davis MIND Institute), the project is a National Institutes of Health Autism Centers of Excellence (ACE) award.

The Vanderbilt site is led by Paul Yoder, Ph.D., professor of Special Education, with co-investigator Zachary Warren, Ph.D., associate professor of Pediatrics and TRIAD director.

The two styles of early intervention being compared are a play-centered model based on the Early Start Denver Model and the discrete trial focused model, both of which have shown effectiveness.

"This research could have important implications for families and health care systems seeking intensive and therefore expensive but needed treatment for children with autism," Yoder said.

The Early Start Denver Model, which was developed by Rogers and colleagues, is a comprehensive behavioral early intervention approach for children with autism, ages 12 to 48 months. The program encompasses a developmental curriculum that defines the skills to be taught at any given time and a set of teaching procedures used to deliver this content. Its core features include naturalistic applied behavioral analytic strategies, sensitivity to typical developmental sequence, deep The discrete trial focused model is based on work from Ron Leaf, John McEachin, and colleagues from the Autism Partnership program in California. Discrete trial training is a one-to-one instructional approach used to teach skills in a planned, controlled, and systematic manner. Each trial or opportunity has a definite beginning and end, with



parental involvement, focus on interpersonal exchange and positive emotional expression, shared engagements with joint activities, and language and communication taught inside a positive, affect-based relationship. praise and/or tangible rewards used to reinforce learning. Early intensive behavioral interventions focused on discrete trial teaching have been used across many early intervention programs for the past two decades and have been highlighted in research reviews as some of the most powerful for preschool children with autism.

The Vanderbilt intervention study will be conducted in Clarksville, Tennessee, with Army families based at Fort Campbell. Children under 2 years of age will be randomly assigned to one of the two interventions and randomly assigned to receive either 15 hrs/week or 25 hrs/wk of in-home therapy over a year. In both interventions, parents will receive 1.5-2 hour training sessions, 2 per month. Parent-child interaction at home and a 24-hour child vocal record

will be recorded monthly. Children's developmental progress will be assessed monthly at Vanderbilt. Therapists will be supervised by Board-Certified Behavioral Analysts.

VANDERBILT KENNEDY CENTER

Remembering Nancy Huffman

N ancy Huffman, longtime VKC Coordinator of Grants Management and Fiscal Services, passed away on April 29. She was 61. VKC staff and faculty who worked with her for the last 36 years remember her expertise in grant budget development and grants management, as well as her warm smile and gentle personality. Co-worker Laura McLeod wrote, "If I ever needed something, even with all she had going on, I could walk into her office and know that she would greet me pleasantly and help me without complaint. People always felt comfortable approaching her. That's a pretty powerful thing, as simple as it sounds. She was, quite simply, one of the kindest people I have ever met in my entire life."

Remembering Bob Newbrough



. Robert (Bob)

Newbrough, Ph.D., a founder of the field of community psychology, died January 2. He was 78. Newbrough, professor of Psychology, emeritus, directed the Center for Community Studies (1966-1980), an early research unit of the Kennedy Center, and later served as the Kennedy Center's associate director (1981-1983). Newbrough's contributions to community psychology earned him many accolades. He was named fellow the American Psychological Association's Division of Community Psychology (1972), was elected Division president (1978), received the Division's award for Distinguished Contributions to Community Research and Theory (1994), and edited the Journal of Community Psychology (1974-1988). The Inter-American Society of Psychology honored him (1989). He co-founded Peabody's innovative Transactional Ecological Psychology Doctoral Training Program in the early 1970s and directed it until 1986. He established and directed Peabody's nationally known doctoral program in Community Psychology. During the late 1990s, he developed and directed the doctoral program in Community Research and Action in Human and Organizational Development.

Friend and colleague Dee Newman wrote, "what distinguished [Bob] . . ., beyond his intellect, was not just his courage and integrity, his kindness and generosity, . . ., it was simply his concern and willingness to listen and offer help and compassion to others."



Leading the Vanguard of Discovery

Zachary Warren, Ph.D.

Associate Professor of Pediatrics, Psychiatry, and Special Education • Director of Treatment and Research Institute for Autism Spectrum Disorders (TRIAD) • Director of Autism Clinical Services, Division of Developmental Medicine, Pediatrics • Vanderbilt Kennedy Center Investigator • Joined Vanderbilt Kennedy Center 2006

Research Interests

My research interests focus on enhanced early detection and intervention for young children as well as the development of technologies to enhance the lives of individuals with autism. I am working on projects that study infants at high-risk for autism in hopes of developing methods for earliest detection, which early behavioral intervention programs work best for specific children, as well as training programs for building system capacity for early accurate diagnosis and effective treatment for families in their own backyards.

Investigator

- Adaptive Response Technology for Autism Spectrum Disorders Intervention, National Institute for Mental Health
- Autism Treatment Network: Cooperative Multi-Center Program for Research and Treatment of Autism, Autism Speaks
- Early Detection of Developmental Disorders, National Institute of Child Health and Human Development

Clinical Interests

My clinical interests revolve around improving systems of care for individuals with autism and their families, especially improving early identification and intervention. Unfortunately, the process of developing initial concerns, discussing concerns, obtaining accurate diagnosis, and translating this information into effective treatment remains a complex, distressful one for most families. I believe there is a great opportunity to improve this challenging context and to link families, providers, and systems of care under the shared mission of acting early and effectively to improve the lives of children with autism.

Education

- B.A., 1998, Psychology and Philosophy, The College of William and Mary
- M.S., 2002, Clinical Psychology, University of Miami
- Ph.D., 2005, Clinical Psychology,

University of Miami

- Clinical Internship, 2004-2005, Children's Hospital Boston/Harvard Medical School
- Postdoctoral Fellowship, 2005-2006, Medical University of South Carolina

Attraction to Developmental Disabilities Research

I spent most of my graduate training learning about the development of very young children who had experienced serious life stressors. This focus on early social and emotional development was matched with opportunities to work with young children with autism during my fellowship. Since then, I have worked with young families concerned that their child may have autism. It is a challenging and powerful experience to try to provide families with answers to questions regarding autism and other developmental concerns. In my opinion, this represents an opportunity to understand how we can exceptionally join with families to create systems of care of meaning and value.

Reasons for VKC Membership

I have been fortunate in that I have been actively involved with the Kennedy Center since arriving at Vanderbilt some 7 years ago. My continued attraction to the VKC is linked to the Center's ability to help form partnerships among researchers, clinicians, educators, and families in a manner that truly impacts communities. The ability to collaborate and work alongside leading scientific experts in behavior, education, genetics, and neuroscience who share a focus on pushing science of impact and meaning is so very fulfilling. I also am fortunate in that increasingly I have been able to intersect with the powerful training programs (i.e., LEND, UCEDD, TRIAD) supported through our Center that provide exceptional training to our next generation of scientific and clinical leaders. In my experience, few places are capable of bringing together current and future leaders in the field of developmental disabilities in such powerful ways.

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Accolades by elizabeth turner



Erik Carter, Ph.D., associate professor of Special Education, received the **2012** Young Professional Award at the AUCD Conference Dec. 2012, in recognition of his "advocacy, commitment, and promise." His work was described as having "an extraordinary impact on

Erik Carter, Ph.D.

preparing youth and young adults with severe disabilities for self-determined lives with meaningful work and strong community support."

David Dickinson, Ed.D., professor of Education, is serving on the Early Literacy Expert Roundtable Panel for the National Governors Association Center for Best Practices.

Carolyn Hughes, Ph.D., professor emerita of Special Education, and Erik Carter, Ph.D., associate professor of Special Education, published The New Transition Handbook: Strategies High School Teachers Use That Work! (Baltimore: Paul H. Brookes Publishing), which provides more than 500 researcher-based, teacher-tested support strategies.

Evon Batey Lee, Ph.D., associate professor of Pediatrics, Psychology, and Psychiatry, was named director of Training for the Vanderbilt Kennedy Center for Excellence in Developmental Disabilities. She succeeds Terri Urbano, Ph.D., professor emerita of Clinical Pediatrics, who retired in December. Lee was also named associate director of Vanderbilt LEND Training.

Autism Speaks has released the ATN/AIR-P's "Quick Tips for Improving Sleep for Children With Autism," created by Beth Malow, M.D., M.S., Burry Chair in Cognitive Childhood Development, professor of Neurology and Pediatrics, and director, Vanderbilt Sleep Disorders Division.

The Tennessee Supreme Court launched an online advocacy video for the public and lawyers as part of its Access to Justice Initiative. Providing Legal Services to Persons With Disabilities was developed by the Disability Law & Advocacy Center of TN and the VKC, and features Elise McMillan, J.D., VKC UCEDD co-director and senior associate in

Psychiatry.



Velma McBride Murry, Ph.D., Betts Chair and professor of Education and Human Development, was recently honored with the Society for Research in Child Development's 2013 **Distinguished Contributions**

Velma McBride Murry, Ph.D.

to Cultural and Contextual Factors in Child Development Award. The award

recognizes an SRCD member's lifetime contributions to the role of culture and context in the understanding of child development.

Tyler Reimschisel, M.D., assistant professor of Pediatrics and Neurology and director, Division of Developmental Medicine and LEND Training, was recently named Vice Chair for Education, Department of Pediatrics.



Courtney Evans Taylor, M.Div., VKC associate director of Communications and Dissemination, received the Integrity Values Award of The Arc Tennessee for her positive impact in the lives of people with intellectual and developmental disabilities.

Courtney Evans Taylor, M.Div.

Jeremy Veenstra-Vander

Weele, M.D., assistant professor of Psychiatry, Pediatrics, and Pharmacology, was among researchers invited to take part in the press conference of the May International Meeting for

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Selected Publications

- Warren, Z. E., Stone, W. L., & Humberd, Q. (2009). A training model for the diagnosis of autism in community pediatric practice. Journal of Developmental and Behavioral Pediatrics, 30, 442-446
- Warren Z. E., McPheeters, M. L., Sathe, N. A., Foss-Feig, J. H., Glasser, A. M., Veenstra-VanderWeele, J. (2011). A systematic review of early intensive intervention for autism spectrum disorders. Pediatrics, 127, e1303 -e1311.

Taylor, J. T., & Warren, Z. E. (2012). Maternal depressive

symptoms following autism spectrum diagnosis. Journal of Autism and Developmental Disorders, 42, 141-1408. Warren, Z. E., Foss-Feig, J., Malesa, E., Lee, E. B., Taylor, J. L.,

- Davidson, J., Newsom, C., & Stone, W. S. (2012). Neurocognitive and behavioral outcomes of vounger siblings
- of children with autism spectrum disorder at age five. Journal of Autism and Developmental Disorders, 42, 409-18.
- Bekele, E., Crittendon, J., Swanson, A., Sarkar, N., & Warren, Z. E. (in press). Pilot clinical application of an adaptive robotic system for young children with autism. Autism.

Autism Research (IMFAR) in Spain, where he highlighted his study "Promising Results on Arbaclofen for Treating Autism Social Withdrawal."

Two VKC investigators received awards at the annual meeting of the American Association on Intellectual and Developmental Disabilities



(AAIDD). Zachary Warren, Ph.D., associate professor of Pediatrics and Psychiatry, received the AAIDD 2013 Early Career Award. Paul Yoder, Ph.D., professor of Special Education, received the AAIDD 2013 Research Award.

Paul Yoder, Ph.D.

Several VKC researchers have

been honored with endowed chairs. Ariel Deutch, Ph.D., professor of Psychiatry and Pharmacology and director, National Parkinson Foundation Center of Excellence, was named the James G. Blakemore Chair in Psychiatry. Isabel Gauthier, Ph.D., professor of Radiology & Radiological Sciences and Psychology, was named the David K. Wilson Chair. Paul Newhouse, M.D., professor of Psychiatry, Pharmacology, and Medicine and director of the Center for Cognitive Medicine, was named the Jim Turner Chair in Cognitive Disorders. R. Jay Turner, Ph.D., professor of Sociology, was named the Harvie Branscomb Chair.

The research of multiple VKC investigators was cited in the update of the Interagency Autism Coordinating Committee Strategic Plan for Autism Spectrum Disorder Research, U.S. Health and Human Services. Included was the research of Suzanne Goldman, Ph.D. (Neurology, Sleep Division); Beth Malow, M.D. (Neurology); Melissa McPheeters, Ph.D. (Obstetrics & Gynecology); James Sutcliffe, Ph.D. (Molecular Physiology & Biophysics), Julie Lounds Taylor, Ph.D. (Pediatrics); Lily Wang, Ph.D. (Biostatistics); Zachary Warren, Ph.D. (Pediatrics); Jeremy Veenstra-VanderWeele, M.D. (Psychiatry); and Paul Yoder, Ph.D. (Special Education). Malow, Warren, and VKC director Elisabeth Dykens, Ph.D. (Psychology), were part of the external strategic planning group for the federal report. Dykens said the extensive citing of Vanderbilt research reflects "Vanderbilt's leadership role in helping our nation answer the urgent questions 'what do we know?' and 'what do we need?' across the lifespan for the growing number of individuals on the autism spectrum."

New LEND Leadership by JAN ROSEMERGY

he new year began with new leadership for the Vanderbilt LEND Training Program, with the January appointment of Tyler Reimschisel, M.D., as director and, in February, Evon Lee, Ph.D., as

associate director. "Tvler was chosen for his deep commitment to interdisciplinary training in developmental disabilities and his experience with the LEND as associate director since 2008," said



Tyler Reimschisel, M.D.

Reimschisel is assistant professor

Elisabeth Dykens,

VKC director.

of Pediatrics and Neurology, director of the Division of Developmental Medicine and the Center for Child Development, associate director of the Pediatric Residency Program, and vice chair for Education, Department of Pediatrics.

Reimschisel succeeded LEND director Terri Urbano, Ph.D., M.P.H., R.N., professor of Clinical Pediatrics, emerita.

"I want to express our deep appreciation to Dr. Urbano for her outstanding leadership as director since LEND's affiliation with the Vanderbilt

of life for children and adults with developmental disabilities and their families." 'Tyler has been a remarkable partner in our evolving LEND program," said retiring LEND

Kennedy Center in 2008," Dykens said. "She has

been at the forefront of national leaders in teaching,

training, and advocacy to improve health and quality

director Terri Urbano. "I'm excited about the new vision he will bring to take LEND to the next level." Evon Batey Lee, Ph.D., who was appointed VKC

UCEDD director of training in January, succeeding Urbano, will now serve as LEND associate director. Lee is associate

professor of Pediatrics

"Evon was chosen

for both her LEND

because of her career-

long commitment to

(clinical-educator),

Psychology, and

Psychiatry.

and UCEDD

leadership roles

mentoring pre-

continuing

professionals and



Evon Batey Lee, Ph.D.

professionals in the wide array of fields involved in treating and caring for children and adults with developmental disabilities and their families," Dykens said.

Lee has held a variety of roles within the VKC and Vanderbilt Pediatrics since she completed a doctorate in psychology at Vanderbilt in developmental and clinical psychology in 1980. Currently, she is VKC coordinator of Psychological Assessment.

The mission of the Vanderbilt LEND is to reduce and prevent neurodevelopmental and related disabilities in children and youth and to increase access to family-centered, community-based, culturally competent, interdisciplinary services. The Vanderbilt LEND is federally funded by the Maternal and Child Health Bureau, U.S. Department of Health and Human Services.

The LEND program focuses on preparing health professionals to assume leadership roles and to develop interdisciplinary team skills, advanced clinical skills, and research skills, in order to meet the complex needs of children with neurodevelopmental and related disabilities. The Vanderbilt LEND includes faculty from Vanderbilt University, Belmont University, Tennessee State University, Meharry Medical College, and the University of Tennessee, as well as affiliates from Family Voices of Tennessee. During the 2012-13 academic year, the LEND had 20 trainees in deaf education, developmental/behavioral pediatrics, nursing, occupational therapy, pediatric audiology, physical therapy, psychiatry, psychology, social work, speechlanguage pathology, and a family trainee.

LEND and Psychiatry—A Good Match BY ELIZABETH TURNER

Danica Denton, M.D.

anderbilt LEND trainees typically come with backgrounds in audiology, occupational and physical therapy, and nursing, among other fields,

but this year, LEND welcomed its first trainee in psychiatry.

Danica Denton was inspired at a young age by her father, a therapist who always wanted to go to medical school. She found her niche in psychology at the University of Arizona and pursued medical school after working alongside psychiatrists as a case manager at a community mental health agency in Tucson.

"I tried to keep an open mind during my medical school rotations and not pigeonhole myself to

psychiatry," Denton said. "However, when I asked my husband if he thought I should go for surgery or pediatrics, he replied that my eyes always lit up when I talked about my psychiatry rotations and that I was much happier there than on any other service."

Denton was matched with Vanderbilt for her

residency. "My first experience with children came on my very first rotation as an intern on the Child and Adolescent Inpatient Unit at the Vanderbilt Psychiatric Hospital, where I had two patients with autism. I really enjoyed working with them, and I realized that it took a team approach to care for these patients."

After indicating her interest in autism to Psychiatry Department chair Stephan Heckers, M.D., Denton was put in contact with

VKC autism researchers in Psychiatry, Blythe Corbett, Ph.D. and Jeremy Veenstra-VanderWeele, M.D.

"While I was planning my fourth year of residency, Dr. Veenstra-VanderWeele described LEND to me. I was immediately fascinated because I had felt for some time that a multidisciplinary approach in the treatment of children and adolescents with ASD was important," she said. "LEND does a great job of allowing everyone who has an interest in ASD to show how their experience can complement everyone else's to benefit patients."

Denton also is working on autism research projects with Veenstra-VanderWeele. One study involves the use of the Autism Treatment Network (ATN) database to look at differences between children who are and are not prescribed atypical antipsychotics.

"I have learned a huge amount from others in LEND. I have learned how each discipline contributes to the care of children with ASD and how what each of us contributes can complement all the other pieces," Denton said.

VKC UCEDD Reaching Out BY COURTNEY TAYLOR

(1) Proud Graduates

Next Steps at Vanderbilt graduated its third class of students on April 24. Carrie DePauw, Will McMillan, and Matthew Moore completed this 2year certificate program for students with intellectual disabilities, which aims to develop academic, social, and independent living skills and to broaden career options. Family, friends, Ambassadores, and Vanderbilt faculty and staff were in attendance to celebrate what Cynthia Cyrus, associate provost for Undergraduate Education, coined "cheerful change." "We come together with joy in what you have accomplished," Cyrus said. "We take pride in the person you've become, and we celebrate now in anticipation of what you are yet to do."

(2) Advocating on "The Hill"

Advocacy activities abound. The VKC and The Arc Tennessee collaborated to produce a booklet for legislators sharing the stories of families on the waiting list for home- and community-based waiver services. VKC UCEDD and LEND trainees and Next Steps students joined several hundred others for Disability Day on the Hill and Vanderbilt Day on the Hill on January 30 at Legislative Plaza. VKC Community Advisory Council members attended the Disability Policy Seminar in Washington, D.C., in mid-April and visited Tennessee legislators.

(3) Education Advocates for Families

The Volunteer Advocacy Project (VAP) graduated its 10th group of advocates on April 15. Across multiple Tennessee sites during 11 weekly sessions, the project trains volunteers in special education law and advocacy strategies. In turn, these trained volunteers assist parents of children with disabilities as they advocate for educational supports. To date, the VAP has trained 293 advocates, and 350-400 families have been referred through The Arc and STEP.

(4) Showing What's Possible!

Two public awareness videos have been produced through the VKC. *Im Thinking College, Even With My Disability* showcases the experiences of four students enrolled in Next Steps at Vanderbilt and was a collaborative project with the Tennessee Alliance for Postsecondary Opportunities for Students With Intellectual Disabilities. *Connecting to the Future: Employment and Disabilities* was developed in collaboration with the Tennessee Developmental Disabilities Network. It highlights two advocates who have jobs based on their own personal interests and the steps they took to get those positions. Both videos were created by Kyle Jonas and can be viewed at kc.vanderbilt.edu.

(5) NOLA Community Service

"As a senior, I get to experience a whole lot of new things. One of which is how I went to New Orleans for the Best Buddies Spring Break. The things we saw were amazing. We saw buildings that were destroyed by Katrina. Katrina was one of the worst tragedies that happened to New Orleans. It was hard to see them, but it reminded me of what happened to us here in Nashville during the flood. While we were there, we helped a church by cooking and building bookshelves, but I like cooking more! I got to hang out with all of the Best Buddies. I had a really good time. Sometimes we got lost, but it was ok!" - Will McMillan, Next Steps at Vanderbilt Class of 2013



Grand Ole Heart Pete Fisher BY COURTNEY TAYLOR

Pete Fisher, Vice President and General Manager of the Grand Ole Opry, remembers the first year the ACM Lifting Lives Music campers performed on the Opry stage. After the performance, he received a phone call from the manager on duty, who told him that a couple and

their daughter wanted to speak with him. The family was visiting Nashville and had come to the Opry not knowing the campers would be performing. He learned that their daughter had Williams syndrome, and she had never before met another person with Williams syndrome. "They were

beyond excited," said Fisher. "We just had



Pete Fisher

to bring them backstage and, of course, all the campers hugged her. It really blindsided us. I'd be lying if I said it wasn't a leap of faith to have them on the stage, but when we saw that first performance and when that family came forward, it was total and complete validation that we should be doing this. We have extraordinary moments every single week at the show, and when the campers are here it's one of the highlights of the year."

Fisher is no stranger to celebrating the skills and talents of people with disabilities. His wife of

> 25 years, Hope, has a sister with Down syndrome. He has witnessed first-hand some of the challenges that she and the family have had finding supports and garnering greater understanding in the community. Fisher and his wife also have two boys, Sean, who is 24 and lives in Los Angeles, and 18-year-old Chris, who will be heading off to college in August.

"Both of our boys are drummers," said Fisher. "They have received very powerful experiences and benefits from music education. I've been involved in an annual fundraiser at their high school called Music 4

Music. It just stems, quite honestly, from an indebtedness that I feel to music education. Music is powerful. All of us here at the Opry have a passion for music and for putting

smiles on people's faces. I think the campers help us develop an even deeper appreciation. Upon observing them and seeing the love they have for music and their experience on the Opry, you can't help but look inward. You can't help but be reminded that music is a gift and that the Opry is a really special place."

Fisher is a graduate of the Leadership Music Program, which serves as a model for the new ACM Lifting Lives Series at Vanderbilt. The 9session, 8-month series gives participants with disabilities an opportunity to learn about different aspects of the music industry, while providing opportunities for enhancing vocational and social skills. Fisher serves on the board of the Lifting Lives Foundation and is excited about the new series, especially since he believes it will expand and extend the benefits of the week-long camp experience.

"The Nashville music industry is extremely giving of its time, its resources, and its people," said Fisher. "You know, there was a time when I thought I had limited capacity, but when I realized my capacity was much larger than I thought it was, it was transformative. Have you ever noticed that the most fulfilled and happiest people are the people who give of their time and resources? Giving back fulfills me more than anything else I do. It's a cliché, but you hear it all the time that it's better to give than receive. And it is! I truly believe that if you want to realize your greatest potential, it has nothing to do with receiving. It has everything to do with giving."

Aging and Cognition in Down Syndrome Hobbs Discovery Grant BY JAN ROSEMERGY

A lthough Down syndrome is the most common genetic cause of intellectual disability, it has received less research attention than autism and other less common disorders. Thanks to advances in medical care, most especially corrective surgery for heart defects, life expectancy for individuals with Down syndrome has increased to 50 to 60 years. Unfortunately, over 50% of adults with Down syndrome develop dementia by age 60. This makes age-related dementia in older adults with Down syndrome an urgent public health concern.

Paul Newhouse, M.D., known nationally for his research on aging and cognition, is now exploring an innovative therapeutic approach for dementia in Down syndrome, thanks to a VKC Nicholas Hobbs Discovery Grant. Newhouse is Jim Turner Professor of Cognitive Disorders and professor of Psychiatry, Pharmacology, and Medicine.

Newhouse was recruited to Vanderbilt to direct the Center for Cognitive Medicine in the

Department of Psychiatry. The Center for Cognitive Medicine conducts research and clinical activities that focus on disorders of cognition (memory, thinking, and learning). A major focus is on the aging brain. The Center also studies latelife cognitive disorders, including Mild Cognitive Impairment (MCI) and Alzheimer disease. MCI is the stage when others notice that an individual is developing mild memory or thinking problems. Many persons with MCI go on to develop Alzheimer disease.

In 2012, Newhouse published study findings in Neurology that suggested wearing a nicotine patch may help improve memory loss in older adults with mild cognitive impairment. Nicotine stimulates cholinergic receptors in the brain that are important for thinking and memory, and it may have neuroprotective effects. People with Alzheimer disease lose some of these cholinergic receptors. Newhouse and VKC director Elisabeth Dykens, Ph.D., are conducting a memory treatment research study for adults with Down syndrome. The study will examine the treatment of memory using lowdose nicotine patches.

Earlier research has shown that brain changes characteristic of Alzheimer disease occur prior to symptoms of memory loss. This suggests that for treatments to be effective, they must begin early. This study will focus on adults with Down syndrome ages 35 and over who are non-smokers and have only mild changes in memory, behavior, and functional independence. The study will involve 6 visits to evaluate medical status, level of functioning, cognitive status, and changes in cognitive ability, memory, attention, and brain wave activity. Treatment will be a 1-month trial of lowdose, FDA-approved nicotine patches. Interested families may contact (615) 322-2082, asante.kamkwalala@vanderbilt.edu.

Honoring Hobbs Society Members

The Vanderbilt Kennedy Center and its Leadership Council thanked members of the Nicholas Hobbs Donor Society at a reception on April 23 at the home of Sue and Andy Spickard. The generosity of Hobbs Society donors supports innovative research on the causes and treatments of disabilities such as autism, Down syndrome, and other genetic syndromes.

Dr. Bob Dittus, associate vice chancellor for Public Health and Health Care at Vanderbilt Medical Center, described the Kennedy Center as a "beacon of hope." He expressed gratitude to Hobbs Society members whose philanthropy is vital to the Center's mission of facilitating discoveries and making positive differences in the lives of persons with developmental disabilities and their families.

VKC director Elisabeth Dykens thanked members whose gifts support Hobbs Discovery Grants. Across types of disabilities and across the life span, she gave several examples of how promising findings from these innovative "seed" grants are a good investment.

Donna Eskind, Leadership Council chair, closed the evening's program, thanking Hobbs members for their important work.

(1) Andy and Sue Spickard, hosts (2) Donna and Jeff Eskind and Elisabeth Dykens (3) Annette Eskind, Elise McMillan, William Anderson Spickard III, and Cathy Brown (4) Elisabeth Dykens, Robert Dittus, Annette Eskind (5) Mary Carlson, Stephen Pert, Beth Malow, and Karen Morgan. Photos by Tommy Lawson.











Grants Awarded

Genetic Analysis of Synapse

Formation and Function Kendal Broadie, Ph.D. (Biological Sciences) National Institute of Mental Health

CamKII Endocannabinoids, Synaptic Plasticity and Motor Function

Roger Colbran, Ph.D. (Molecular Physiology & Biophysics) National Institute of Neurological Disorders and Stroke

Word Problems, Language, and Comorbid Learning Disabilities Lynn Fuchs, Ph.D. (Special Education) National Institute of Child Health and Human Development

Development of a Manualized Wireless Moisture Alarm Intervention for Toilet Training Children With Autism Whitney Loring, Psy.D. (Pediatrics) Organization of Autism Research

Functional Mapping of Cortical Networks in Primates With Laser Stimulation Anna Roe, Ph.D. (Psychology) National Institute of Mental Health

Sox 10 Alleles for Functional Analysis of Glial Lineages Michelle Southard-Smith, Ph.D. (Medicine) National Institute of Neurological Disorders and Stroke

Study of Oxytocin in Autism to Improve Reciprocal Social Behaviors Jeremy Veenstra-VanderWeele, M.D. (Psychiatry) National Institute of Child Health and Human Development

Translating OCD Gene-Association Studies Into Mice to Examine SLC1A1 Function Jeremy Veenstra-VanderWeele, M.D. (Psychiatry) National Institute of Mental Health

VKC Nicholas Hobbs Discovery Grants

Treatment Effects on Face Processing in Autism Blythe Corbett, Ph.D. (Psychiatry)

Oxytocin and Multisensory Integration in the Neonate Elizabeth Hammock, Ph.D. (Pediatric Endocrinology)

Randomized Clinical Trial of Pacifier Activated Lullaby (PAL) Nathalie Maitre, Ph.D., M.D. (Neonatology/Pediatrics)

Linking Vanderbilt Autism Research Registry to State Databases Richard Urbano, Ph.D. (Pediatrics)

Adaptive Robotic Intervention Architecture for Autism Spectrum Disorders Zachary Warren, Ph.D. (Pediatrics)

Nicotinic Treatment of Age-Related Cognitive Decline in Down Syndrome: A Pilot Trial Paul Newhouse, M.D. (Psychiatry)



Leadership Council of Vanderbilt Kennedy Center

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Breakthroughs in Developmental Disabilities Research



View a brief video of 11 national experts in developmental disabilities research reflecting on the most exciting breakthroughs of the last decade and what holds the greatest promise for the future.

kc.vanderbilt.edu/ DDbreakthroughs

Video produced by Blythe Corbett, Ph.D. Tony Maupin, videographer Discovery is a quarterly publication of the Vanderbilt Kennedy Center designed to educate our friends and the community, from Nashville to the nation. The Center is a *Eunice Kennedy Shriver* Intellectual and Developmental Disabilities Research Center funded by the *Eunice Kennedy Shriver* National Institute of Child Health and Human Development, and a University Center for Excellence in Developmental Disabilities (UCEDD) funded by the Administration on Intellectual and Developmental Disabilities (AIDD). *Discovery* is supported in part by Grant No. HD 15052 from EKS NICHD, AIDD Grant #90DD0595, and LEND Training Grant No. T73MC00050 MCHB/HRSA.

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Elisabeth Dykens, Ph.D., Kennedy Center Director; Karoly Mirnics, M.D., Ph.D., Associate Director; Jan Rosemergy, Ph.D., Deputy Director and Director of Communications; Tim Stafford, Director of Operations

UCEDD: Elisabeth Dykens, Ph.D., Co-Director; Elise McMillan, J.D., Co-Director; Evon Lee, Ph.D., Training; Robert Hodapp, Ph.D., Research; Jan Rosemergy, Ph.D., Dissemination

LEND: Tyler Reimschisel, M.D., *Director*; Evon Lee, Ph.D., *Associate Director*

TRIAD: Zachary Warren, Ph.D., Director

Discovery: Editor: Jan Rosemergy, Ph.D.; Graphic Designer: Kylie Beck; Photo Editor: Amy Pottier

Photographers: VKC: Kylie Beck, Tammy Day, Tony Maupin, & Elise McMillan; Vanderbilt: Dan Loftin, Anne Rayner, John Russell, & Susan Urmy Other: Tommy Lawson

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CALENDAR OF EVENTS | JUNE-SEPTEMBER 2013



Unless otherwise noted, events are free and open to the public. Events are subject to change. Please check the website calendar at kc.vanderbilt.edu or contact (615) 322-8240 or toll-free (1-866) 936-VUKC [8852].

Please keep this calendar and check the Event Calendar on the VKC website for updates. If you wish to receive event announcements by email, send your email address to kc@vanderbilt.edu.

For disability-related training and other events statewide and nationally, see the searchable Pathfinder Disability Calendar www.familypathfinder.org.

*Event will be held in Room 241 Vanderbilt Kennedy Center/One Magnolia Circle Bldg (110 Magnolia Circle).

**VKC Member or Investigator

JUNE 22*

TRIAD Families First Workshops Developing Successful Sleep and Feeding Habits Register at kc.vanderbilt.edu/ registration. Info (615) 322-6027 or families.first@vanderbilt.edu Saturday 9 a.m.-12 p.m.

■ JUNE 25-26*

Autism Diagnostic Observation Schedule (ADOS-2) Clinical Training

TRIAD workshop for psychologists, pediatricians, behavioral specialists, and SLPs. Fee: \$300. Register at kc.vanderbilt.edu/registration. Info amy.r.swanson@vanderbilt.edu or (615) 322-6533. Tuesday-Wednesday 8 a.m.-4:30 p.m.

JUNE 29*

Clinical Genetics for Health Care Professionals

Workshop by Vanderbilt Pediatrics Developmental Medicine Division for health care professionals who are not geneticists. \$50 (includes lunch) 7.25 CME & APA CE. Register at kc.vanderbilt.edu/registration. Info pam.grau@vanderbilt.edu Saturday 7:30 a.m.-2 p.m.

JULY 20*

TRIAD Families First Workshops Building Functional Social and Play Skills Register at kc.vanderbilt.edu/ registration. Info (615) 322-6027 or families.first@vanderbilt.edu Saturday 9 a.m.-12 p.m.

JULY 22*

Statistics and Core Methodology Training Seminar Improving Research Using Advanced REDCap Interfaces Scott Burns, Neuroimaging Analyst Register at kc.vanderbilt.edu/registration Monday 12:30-1:30 p.m.

JULY 30-31

School Speech-Language Pathology Conference Two lunch-time plenary sessions and selection among 6 (90-minute) small group sessions (1.1 ASHA CEUs). Register at kc.vanderbilt.edu/registration Info languagelab@vanderbilt.edu 8th Flr Lecture Hall, Vanderbilt Bill Wilkerson Center, Medical Center East Tuesday-Wednesday 8 a.m.-3 p.m. **Team William 2013**. Despite pouring rain, Andrea McDermott Sanders (VU Peabody M.Ed. '08) ran the Music City Marathon to continue annual Team William fundraising to endow VKC Reading Clinic scholarships for students with Down syndrome and to fund research on literacy in Down syndrome. Photo: Margaret and Dr. William Anderson Spickard III, Sanders, and William Spickard.

AUGUST 17*

TRIAD Families First Workshops Addressing Challenging Behaviors Register at kc.vanderbilt.edu/ registration. Info (615) 322-6027 or families.first@vanderbilt.edu Saturday 9 a.m.-12 p.m.

SEPTEMBER 13*

Community Advisory Council Meeting Info (615) 936-8852 Friday 9 a.m.-2 p.m.

SEPTEMBER 18

Neuroscience Graduate Program Seminar Series TBA

Stephen J. Smith, Ph.D., Professor of Molecular & Cellular Physiology Stanford University. Co-sponsor Vanderbilt Brain Institute Room 1220 MRB III Lecture Hall Wednesday 4:10 p.m.

SEPTEMBER 26*

Lectures on Development and Developmental Disabilities Emotion and Survival: What's the Relation? Joseph LeDoux, Ph.D., Professor of Neural Science & Psychology, New York University Thursday 4:10 p.m.



CALENDAR OF EVENTS | JUNE-SEPTEMBER 2013

VKC SUMMER PROGRAMS

Info laura.mcleod@vanderbilt.edu *Registration is closed. Contact TN Disability Pathfinder for information on summer activities.

• JUNE 3-15

SENSE Theatre Camp* For youth, 7-18 years of age, with and without autism spectrum disorders. Performances Friday, June 14 and Saturday, June 15, 7 p.m. at University School of Nashville Auditorium, 2000 Edgehill Ave, Nashville, TN 37212. Tickets \$5.

• JUNE 23-29

ACM Lifting Lives Music Camp* Residential camp for individuals with Williams syndrome (16 yrs+). • JULY 14-19

Next Steps at Vanderbilt Summer Institute*. Residential college transition program for rising high school juniors, seniors, and young adults with developmental disabilities up to age 24.

READING CLINIC SUMMER SESSION • JUNE 3-JULY 15*

(no sessions Wk July 4) Tutoring students through middle school. 24 (40-minute) sessions, ranging 8 a.m.-12 noon, M-Th. Info on fall sessions (615) 936-5118 or readingclinic@vanderbilt.edu

ARTS AND DISABILITIES EXHIBIT

Monday-Friday 7:30 a.m.-5:30 p.m. Lobby VKC/One Magnolia Circle Bldg. Info (615) 936-8852

• THROUGH JULY 31 The ART of AUTISM. Artists of Tennessee and Its Neighbors

 AUGUST-SEPTEMBER Sandhill Cranes, Owls, and Elephants. Artists of Pacesetters, Inc.

■ AUTISM TRAININGS FOR K-12 SCHOOL PERSONNEL

With the Tennessee Department of Education, TRIAD offers free autism-specific workshops for school personnel, parents, and the community in locations across the state. For dates/locations, see kc.vanderbilt.edu/TRIAD/events

■ LEARNING ASSESSMENT CLINIC

Multidisciplinary academic assessments of students, 5-25 years, to identify learning strengths and challenges and to recommend strategies to improve academic learning. Info (615) 936-5118 or LAC@vanderbilt.edu

NEXT STEPS AT VANDERBILT

A 2-year certificate postsecondary education program for students

with intellectual disabilities providing individualized Programs of Study in education, social skills, and vocational training. Info (615) 343-0822 or NextSteps@vanderbilt.edu

■ TABS (TENNESSEE ADULT BROTHERS AND SISTERS) NETWORK

TABS Facebook page www.facebook.com/tabs.siblings, info.tabs@vanderbilt.edu

TAKE PART IN RESEARCH

VKC Research Studies For children and adults, with and without disabilities Lynnette Henderson (615) 936-0448

- Toll-free (1-866) 936-VUKC [8852] • Research Family Partners kc.vanderbilt.edu/rfp Register and be notified of research studies
 - StudyFinder kc.vanderbilt.edu/studyfinder View lists of studies, criteria, and contact information
 - See also VUMC Clinical Trials www.vanderbilthealth.com/ clinicaltrials

■ TENNESSEE WORKS

Web hub for information related to employment of people with disabilities. Info (615) 322-4999 TennesseeWorks.org

■ TENNESSEE DISABILITY PATHFINDER ■ MULTI-CULTURAL OUTREACH

Helpline, Web-Searchable Database with Calendar and Resource Library, Print Resources. Project of VKC UCEDD and TN Council on Developmental Disabilities www.familypathfinder.org English (615) 322-8529 Español (615) 479-9568 Toll-free (1-800) 640-INFO [4636] tnpathfinder@vanderbilt.edu

■ VOLUNTEER ADVOCACY PROGRAM

Fall 2013 training (11 weekly sessions) in special education law and advocacy strategies. Info lynise.parisien@vanderbilt.edu

ASMT EVENTS

Autism Society of Middle Tennessee Registration is requested for all events. ASMT members free; nonmembers \$5/family. Vanderbilt Kennedy Center/One Magnolia Circle. Info (615) 385-2077 www.tnautism.org

DSAMT EVENTS

Down Syndrome Association of Middle Tennessee Info (615) 386-9002 www.dsamt.org



Fifth Annual TABS Conference, April 5-6, provided adult brothers and sisters of siblings with disabilities information, networking, and support.

